New H₂ collision-induced absorption and NH₃ opacity and the spectra of the coolest brown dwarfs

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ABSTRACT

We present new cloudy and cloudless model atmospheres for brown dwarfs using recent ab initio calculations of the line list of ammonia (NH_3) and of the collision-induced absorption of molecular hydrogen (H_2). We compare the new synthetic spectra with models based on an earlier description of the H_2 and NH_3 opacities. We find a significant improvement in fitting the nearly complete

spectral energy distribution of the T7p dwarf Gliese 570D and in near infrared color-magnitude diagrams of field brown dwarfs. We apply these new models to the identification of NH_3 absorption in the H band peak of very late T dwarfs and the new Y dwarfs and discuss the observed trend in the $NH_3 - H$ spectral index. The new NH_3 line list also allows a detailed study of the medium resolution spectrum of the T9/T10 dwarf UGPS J072227.51-054031.2 where we identify several specific features caused by NH_3 .

Subject headings: stars: low mass, brown dwarfs — stars: atmospheres

1. Introduction

Brown dwarfs, with masses between $\sim 0.01-0.075\,M_{\odot}$, are not massive enough to maintain steady hydrogen fusion in their cores and, except when they are very young, have effective temperatures below that of main sequence stars. Their low atmospheric temperatures allow the formation of molecules and condensates that leave conspicuous imprints on their spectral energy distribution (SED) that have led to the creation of two spectral classes: the L dwarfs with $T_{\rm eff} \sim 2400-1400\,{\rm K}$ (Kirkpatrick 2005) and the cooler T dwarfs extending down to $T_{\rm eff} \sim 500\,{\rm K}$ (Leggett et al. 2010a). The recent discovery of several extremely cool dwarfs by the Wide-field Infrared Survey Explorer (WISE) motivates the creation of a new spectral class: the Y dwarfs with $T_{\rm eff} \lesssim 500\,{\rm K}$ (Cushing et al. 2011).

The spectra of L dwarfs are characterized by absorption bands of VO, TiO, H₂O, CO, FeH, CrH and the very broad doublets of Na I (0.59 μ m) and K I (0.77 μ m) (Kirkpatrick et al. 1999). While condensates, primarily corundum, magnesium silicates and iron, do not have spectral features as distinctive as those of molecules, their continuum opacity reddens the near-infrared spectrum of L dwarfs significantly. The effects of the cloud opacity peak at $T_{\rm eff} \sim 1500$ –1600 K or spectral type L6–L8. In the cooler T dwarfs, the condensate cloud layer sinks below the photosphere, TiO and VO have been lost to TiO₂, VO₂ and condensates (Burrows & Sharp 1999; Lodders 2002) the carbon chemistry has shifted to favor CH₄ (Burgasser et al. 2002; Lodders & Fegley 2002), and NH₃ bands appear in the mid-infrared around T2 (Cushing et al. 2006). In the very latest T dwarfs and in the new Y dwarfs, weak bands of NH₃ appear in the near-infrared (Delorme et al. 2008; Cushing et al. 2011).

Due to their additional degrees of freedom and greater complexity, the opacities of molecules are not nearly as well understood as those of atoms and ions. The primary molecular opacity data bases are the HITRAN (Rothman et al. 2003, 2005, 2009) and the GEISA

(Jacquinet-Husson et al. 1999, 2005) compilations, which were developed for modeling the Earth's atmosphere and consider the opacity of gases at temperatures of 300 K and below. Since the temperature in brown dwarf atmospheres can be up to 10 times higher, those line lists are missing the so-called "hot bands" that arise from the absorption of a photon from an excited lower level that is not thermally populated at 300 K. In some cases, the molecular line spectrum is so rich that individual transitions are unresolved in laboratory spectra or it becomes impossible to assign the observed transitions to specific molecular energy levels. The line lists are thus incomplete for the conditions encountered in brown dwarf atmospheres, especially for higher photon energies, hot bands, and weak transitions corresponding to near infrared and optical wavelengths. Incomplete molecular line lists limit the predictive capability of atmosphere models and synthetic spectra of brown dwarfs (Cushing et al. 2008; Stephens et al. 2009), a problem that was recognized in the first attempts to model the observed spectrum of a brown dwarf, the T7p dwarf Gliese 229B (Marley et al. 1996; Allard et al. 1996). In view of the difficulties in analyzing laboratory spectra of high temperature molecular gases, an alternative is to compute a molecular line list with a first-principles approach.

The past 15 years of brown dwarf research have motivated the theoretical modeling of the opacity of several key molecules. New molecular line lists for H₂O (Partridge & Schwenke 1997; Barber et al. 2006), CO (Goorvitch 1994), TiO (Schwenke 1998), VO (Plez 1998), FeH (Dulick et al. 2003), CrH (Burrows et al. 2002), have improved the quality of the brown dwarf model spectra significantly. Methane (CH₄) has been and remains the most important molecule observed in brown dwarf spectra (defining the T spectral class, but appearing as early as L5; (Noll et al. 2000)) with a very incomplete line list at near infrared wavelengths although some progress has been made (Freedman et al. 2008; Schwenke 2002). The two most recent developments in this area are new calculations of a line list of ammonia (NH₃) and of the continuum opacity from collision-induced absorption (CIA) by H₂ molecules.

We study the effect of these two much anticipated new opacity calculations on the spectra and colors of brown dwarfs. In section $\S 2$, we discuss the nature of H_2 CIA opacity, how the new calculation changes the opacity and how if affects the SED of cool brown dwarfs. The role of the NH_3 opacity in brown dwarf spectra is presented in section $\S 3$, where we show how the new line list for NH_3 affects primarily the near-infrared spectrum and brings new spectral features that can help define the T/Y dwarf boundary. We demonstrate the resulting improvement in synthetic spectra by comparing with the well-characterized SED of the T7.5 dwarf Gliese 570D and near infrared color-magnitude diagrams in section $\S 4$. The improved NH_3 line list provides the opportunity to comment on the role of ammonia in extremely cool dwarfs near the T/Y transition and on recent detections of this molecule in the spectra of extreme objects (section $\S 5$). Concluding remarks are provided in section $\S 6$.

2. Collision-induced absorption by H_2

Molecular hydrogen H₂ is the most abundant component of brown dwarf atmospheres. Because of its high degree of symmetry, the H₂ molecule has no permanent dipole moment and only quadrupole transitions are allowed, which are very weak. During a collision with another particle, however, the compound H₂-particle system has a temporary dipole moment and can thus undergo a radiative dipole transition. The collision-induced absorption process is significant only at relatively high densities where collisions are frequent, such as in the atmospheres of planets and brown dwarfs. Three-body and higher order collisions also contribute to CIA but are probably negligible in brown dwarf atmospheres (Hare & Welsh 1958).

While the probability of a dipole transition in a typical molecule involves only molecular properties, CIA also depends on the frequency of collisions with other atoms or molecules (primarily other H₂ and He, by far the most abundant species), which is linearly dependent on the density of the gas. For a given metallicity, the relative importance of CIA absorption in a brown dwarf atmosphere is a measure of the density of the gas, which is closely tied to the surface gravity. On the other hand, the importance of H₂ CIA decreases in higher metallicity atmospheres as the absorption from molecules involving heavier elements (CH₄, CO, H₂O, NH₃) becomes dominant. This sensitivity of H₂ CIA to gravity and metallicity is a useful spectral diagnostic in T dwarfs (Burgasser et al. 2002, 2003a; Burgasser 2004b).

The H₂ CIA transitions occur between vibrational-rotational levels of the H₂ molecule and form a series of bands corresponding to $\Delta v = 0$, 1, 2, 3, etc., where v is the vibrational quantum number of the molecule. Each band is formed by transitions between the different rotational levels for a given Δv . Collisional broadening of the individual lines is comparable to the line spacing and thus H₂ CIA is effectively a continuum source of opacity. Borysow and her collaborators calculated the H₂-H₂ and H₂-He CIA coefficients in the temperature regime of interest for brown dwarf and white dwarf atmospheres (Borysow & Frommhold 1986; Borysow et al. 1989; Jorgensen et al. 2000; Borysow et al. 2001; Borysow 2002; Gustafsson & Frommhold 2003)¹, and those are universally used by brown dwarf modelers (Freedman et al. 2008; Sharp & Burrows 2007; Allard et al. 2001). The H₂-H₂ and H₂-He CIA coefficients are shown in Figures 1 and 2, respectively. While these two sources of opacity have similar frequency (ν) and temperature (T) dependences, H₂-He plays a secondary role since the collisions with He are about 10 times less frequent than with H₂ as follows from their relative number densities. The absorption coefficient per

 $^{^1\}mathrm{Additional}$ $\mathrm{H_2}$ CIA references as well as codes and tables can be found at http://www.astro.ku.dk/~aborysow/programs/index.html

H₂ molecule is

$$\kappa_{\rm H_2}(\nu, T) = n_{\rm H_2} \sigma_{\rm H_2 - H_2}(\nu, T) + n_{\rm He} \sigma_{\rm H_2 - He}(\nu, T),$$
(1)

where n_i is the number density of species i and σ_{H_2-i} is the H_2-i CIA cross section. The successive CIA bands corresponding to $\Delta v=0,1,...5$ are clearly visible in Figures 1 and 2. At higher temperatures, weaker transition become more important and fill in the inter-band regions. The $\Delta v=1$ band centered at $\sim 4200\,\mathrm{cm^{-1}}$ (2.4 $\mu\mathrm{m}$) is important in T dwarfs as it coincides with the K band flux peak, which is caused by a minimum in the combined opacities of $\mathrm{H}_2\mathrm{O}$ and CH_4 . On the other hand, the J and H band flux peaks in the spectra of T dwarfs are shaped primarily by $\mathrm{H}_2\mathrm{O}$ and CH_4 absorption. Thus, the K/J and K/H flux ratios reflect the relative importance of H_2 CIA to the $\mathrm{H}_2\mathrm{O}$ and CH_4 opacities which is affected primarily by the gravity and metallicity of the atmosphere. A relatively weaker K band flux indicates a higher gravity (stronger H_2 CIA) or a lower metallicity (weaker $\mathrm{H}_2\mathrm{O}$ and CH_4 bands), or both. Our model atmospheres tend to underestimate the K band flux, resulting in bluer J-K than is observed for late T dwarfs (Saumon & Marley 2008). We have long suspected that this systematic offset comes form an overestimate of the strength of the H_2 CIA in the $\Delta v=1$ band.

2.1. New calculation of the H_2 collision-induced absorption

First-principles calculations of molecular opacities are typically conducted in two steps. First, a potential energy surface is calculated for a range of molecular geometries representing variations in bond lengths and bond angles. This potential surface, which is multi-dimensional for polyatomic molecules, is obtained by solving the quantum mechanical Schrödinger equation for the electrons for each given nuclear geometry using methods of quantum chemistry. The potential surface then serves as input for the calculation of the eigenstates of the molecule and the induced dipole moment surface from which allowed transitions and their oscillator strengths can be calculated to generate a line list. For bands originating from excited states, or for relatively high energy transition (in the optical), the final state is well above the ground state of the molecule. The calculation of these lines requires an accurate knowledge of the potential energy surface far from the minimum that approximately corresponds to the ground state energy, which involves many subtle quantum mechanical effects and requires extensive calculations for widely separated configurations of nuclei. The calculation of higher energy transitions and of "hot bands" is thus increasingly difficult.

The growing evidence from brown dwarf spectra that something was amiss in the $\Delta v = 1$ band of H₂ CIA as well as applications to ultracool white dwarfs (Bergeron, Saumon & Wesemael

1995) prompted a new calculation of the H_2 - H_2 and H_2 -He CIA opacity in the entire regime of temperatures and frequencies of interest (Frommhold et al. 2010; Abel et al. 2011, 2012). New, high accuracy potential energy surfaces and induced dipole surfaces were generated for these two CIA processes. The resulting opacity was calculated on a fine temperature and frequency grid, from $\nu = 20$ to $10^4 \, \mathrm{cm}^{-1}$ and $T = 200 - 3000 \, \mathrm{K}$ for $\mathrm{H_2\text{-}H_2}$ and 20 to $20000 \, \mathrm{cm}^{-1}$ and $T = 200 - 9900 \,\mathrm{K}$ for $\mathrm{H_2\text{-}He}$ (Richard et al. 2012). The $\mathrm{H_2\text{-}H_2}$ table is being extended to the same (T, ν) grid as the H₂-He table. In each case, the entire parameter range is covered in a uniform manner, while earlier tabulations used different approximations and potential surfaces for different temperature and frequency ranges. As is the case for the older calculations, the new H₂-H₂ opacities are in excellent agreement with experimental data, which are limited to $T \leq 300 \,\mathrm{K}$ (Frommhold 1993; Abel et al. 2011). Comparisons with the CIA opacity previously used in modeling brown dwarf atmospheres are shown in Figures 1 and 2. As expected, differences are negligible or minimal at low temperatures. The differences become substantial at 2000 K in the roto-translational band for H_2 - H_2 ($\sim 1000 - 2000 \,\mathrm{cm}^{-1}$) and in all roto-vibrational bands ($\Delta v > 0$). In the fundamental band ($\Delta v = 1$), which affects the K band flux in brown dwarfs, the H_2 - H_2 CIA has decreased by 24% at 1000K and by 44% at 2000K, which goes in the direction required to bring the K band flux of the models into better agreement with the observations. The blue side of the fundamental band of H_2 - H_2 CIA extends into the H photometric band where the absorption has decreased somewhat at temperatures above $\sim 1500 \,\mathrm{K}$. The H₂-He contribution (Fig. 2) shows similar variations, except that the absorption coefficient increases at the peaks of the fundamental and first overtone bands. At frequencies above 10⁴ cm⁻¹, the corrections can be factors of 3 or more. This matters less in brown dwarfs as H₂ CIA is not an important source of opacity at optical wavelengths and the He abundance is $\sim 10\%$ that of H₂.

The effect of the change in the CIA opacity is shown in Fig. 3 where spectra computed with the old and the new opacity but with the same (T,P) atmospheric structures, are compared. All the flux peaks in the near infrared are affected, with rather modest flux increases of $\lesssim 15\%$. Since the new H₂ CIA opacity is nearly identical to the older calculation at $T=300\,\mathrm{K}$ and increasingly differs at higher temperatures, the differences in the spectra tend to be larger at higher T_{eff} and nearly vanish for $T_{\mathrm{eff}}\lesssim 500\,\mathrm{K}$ except in the Y and J bands which are relatively transparent and probe the atmosphere at temperatures well above T_{eff} . Since the CIA opacity increases with the gas density, i.e. with gravity, lower gravity spectra are less affected by the new CIA. For example, at $\log g = 4$ (cgs), the flux in the peaks of the SED increases by less than 7%. At effective temperatures higher than those shown in Fig. 3, a greater change in the spectrum would be expected in cloudless models. However, in brown dwarfs atmospheres with $T_{\mathrm{eff}} \gtrsim 1400\,\mathrm{K}$ the continuum opacity of the condensates overwhelms the H₂ CIA opacity and the spectrum is unaffected.

3. Ammonia opacity

In brown dwarf atmospheres, the chemistry of nitrogen is dominated by molecular nitrogen N_2 and ammonia NH_3 , the latter being favored at higher pressures and lower temperatures (Lodders & Fegley 2002). The transformation of N_2 into NH_3 occurs at $T_{\rm eff} \sim 800~{\rm K}$ and is, along with the similar conversion of CO to CH_4 , one of the most significant chemical changes in brown dwarf atmospheres as $T_{\rm eff}$ decreases. While N_2 has no permanent dipole moment and is effectively invisible in brown dwarf spectra² the fundamental band of NH_3 in the 10–11 μ m region is seen in T2 and later dwarfs (Cushing et al. 2006) and the entire 9–14 μ m region is affected by NH_3 absorption (Saumon et al. 2006). Ammonia has a series of weaker overtone bands in the near infrared, two of which are centered at 2.0 and 1.5 μ m and fall in the K and H photometric bands where the SEDs of T dwarfs show emission. The apparition of these bands in the spectra of very cool brown dwarfs has been suggested as a trigger for a new spectral class (Burrows et al. 2003; Leggett et al. 2007a; Kirkpatrick 2008).

Saumon et al. (2000) reported the detection of three NH₃ features in a R = 3100 spectrum of the T7p Gliese 229B in the $2.02 - 2.045 \,\mu\mathrm{m}$ range but other expected NH₃ features were missing. This tentative detection was followed by a decade where no other near-infrared spectrum of T dwarfs was taken at high enough resolution in the K band to reveal NH₃ fea-The first unambiguous detection of NH₃ in brown dwarfs was achieved with the Infrared Spectrograph of the Spitzer space telescope that clearly showed the 10–11 μ m band in the combined spectrum of ϵ Indi Bab (T1+T6) (Roellig et al. 2004; Mainzer et al. 2007) and then in all dwarfs of type T2 and later observed with the IRS (Cushing et al. 2006; Saumon et al. 2007; Cushing et al. 2008; Stephens et al. 2009; Leggett et al. 2010b). Recently, the search for extremely cool dwarfs has uncovered several objects with $T_{\rm eff} \lesssim 600\,{\rm K}$ where the blue wing of the H band flux peak appears depressed due the presence of NH_3 (Delorme et al. 2008; Lucas et al. 2010; Liu et al. 2011; Cushing et al. 2011). Finally, a new $R \sim 6000$ near-infrared spectrum of the $T_{\rm eff} \sim 500\,\mathrm{K}$ T9 dwarf UGPS J072227.51-054031.2 contains NH₃ features (Bochanski et al. 2011). The detection of NH₃ in the near-infrared spectrum of several ultracool dwarfs is one of the factors that motivate the creation of the new Y spectral class (Cushing et al. 2011).

The analysis of the spectral signature of NH_3 in T dwarfs is complicated by the presence of vertical mixing in the atmosphere and by the invisibility of N_2 . Because the time scale for the conversion of N_2 to NH_3 is very long at low temperatures, gas eddies from very

²Like H₂, N₂ causes collision-induced absorption (Borysow & Frommhold 1986, 1987), but its mole fraction of $\sim 10^{-4.2}$ makes this contribution to the opacity in brown dwarf atmospheres completely negligible.

deep in the atmospheres where N_2 and NH_3 are in chemical equilibrium at high temperatures can be transported vertically by convection and slow mixing in the radiative zone to regions of lower temperatures on a time scale that is much shorter than the $N_2 \rightarrow NH_3$ conversion can take place, resulting in a reduced NH_3 abundance (Fegley & Lodders 1994, 1996; Lodders & Fegley 2002; Saumon et al. 2006; Hubeny & Burrows 2007). This process can reduce the observed abundance of NH_3 by factors of 5-10, weakening the NH_3 bands noticeably. Reduced NH_3 absorption in the $9-14\,\mu\mathrm{m}$ region has been found in every T dwarf observed with the *Spitzer* IRS with a high enough S/N ratio (Saumon et al. 2006; Mainzer et al. 2007; Saumon et al. 2007; Burgasser et al. 2008b; Leggett et al. 2009, 2010b) and can be fully explained by considering the convective mixing time scale and the time scale for $N_2 \rightarrow NH_3$ conversion. Considering the relatively well understood chemistry of nitrogen and the universality of a deep convection zone in the atmospheres of brown dwarfs, it is reasonable to assume that the abundance of NH_3 is reduced by vertical transport in all T dwarfs.

In view of the currently very limited opportunities for spectroscopic observations of brown dwarfs in the mid-infrared and the relative ease of near-infrared spectroscopy, the classification of the growing number of very late T dwarfs and of the new Y dwarfs relies primarily on their near infrared spectral characteristics where NH₃ figures prominently as a new spectral index indicator (Delorme et al. 2008; Lucas et al. 2010; Cushing et al. 2011; Burgasser et al. 2011) as anticipated by Burrows et al. (2003); Leggett et al. (2007a) and Kirkpatrick (2008). The importance of NH₃ as a strong absorber in the mid-infrared, an indicator of non-equilibrium chemistry, and an important marker for the spectral classification of very late T dwarfs and Y dwarfs demands a high quality line list to compute reliable models.

3.1. New theoretical NH₃ line list

Our previous models used a NH₃ line list from the HITRAN data base (Rothman et al. 2003, 2005, 2009) supplemented with room temperature data in the $6000-7000\,\mathrm{cm^{-1}}$ range (Freedman et al. 2008; Sharp & Burrows 2007). In principle, the fundamental band at $10-11\,\mu\mathrm{m}$ should be well represented in the low temperature ($\sim 300\,\mathrm{K}$) HITRAN data base but the line list becomes increasingly incomplete at shorter wavelengths and higher temperatures. The resulting opacity at 400 and 1000 K is shown in Figs. 4 and 5. There are several gaps in the inter band regions $2.5-2.75\,\mu\mathrm{m}$, $\sim 2.11\,\mu\mathrm{m}$ and $1.6-1.88\,\mu\mathrm{m}$, and the list stops at $1.42\,\mu\mathrm{m}$. Such a line list is inadequate to analyze the rapidly growing spectroscopic data for $T_{\rm eff} \lesssim 600\,\mathrm{K}$ dwarfs, which has motivated recent experimental

(Hargreaves, Li & Bernath 2011) and theoretical work on ammonia (Huang, Schwenke & Lee 2011,?; Yurchenko, Barber & Tennyson 2011).

A new extensive line list for NH₃ has recently been calculated from first principles. It is based on a new potential energy surface (Yurchenko et al. 2005) that has been refined by fitting calculated energy levels to nearly 400 levels that are accurately known experimentally (Yurchenko, Barber & Tennyson 2011). The resulting energy levels agree with the experimental values to within $1\,\mathrm{cm}^{-1}$ and generally under $0.1\,\mathrm{cm}^{-1}$. This 'BYTe' NH₃ line list (Yurchenko, Barber & Tennyson 2011) contains over 1.1 billion lines, compared to ~ 34000 in HITRAN. It extends from 0 to $12000\,\mathrm{cm}^{-1}$ (0.83 $\mu\mathrm{m}$) and is applicable up to 1500 K. An independent calculation of a NH₃ line list is under way (Huang, Schwenke & Lee 2011). Figures 4 and 5 show the improvement of the BYTe line list over the HITRAN opacity. At 400 K, we expect the HITRAN opacity to be quite good since it is based on data acquired primarily at room temperature. This is indeed what the BYTe line list shows as both are nearly identical near the center of all the bands in common between the two line lists. In addition, the BYTe line list includes new bands at $1.65 \,\mu\mathrm{m}$ and shortward of $1.4 \,\mu\mathrm{m}$, and contains many weak lines that fill in the inter band regions where there are gaps in the HITRAN line list. The differences become more dramatic at higher temperatures (Fig. 5) where the transitions from thermally excited levels come into play, reducing the contrast between the peaks and valleys of the opacity and increasing the band opacity by factors of $\sim 3-5$ compared to the HITRAN value. The opacity in the $8-15\,\mu\mathrm{m}$ also increases significantly.

The effect of the change in NH₃ opacity on the synthetic spectra is shown in Fig. 6. For this comparison, spectra computed with the HITRAN and the BYTe line lists use the same (T, P) atmosphere structure. At $T_{\text{eff}} = 1200 \,\text{K}$ (not shown), the spectra are identical but at $T_{\rm eff} = 800 \, \rm K$ there is a small flux decrease of $\lesssim 20\%$ on the blue side of both the H and K peaks. At lower T_{eff} , the chemical equilibrium shifts to strongly favor NH₃ over N₂ as the dominant nitrogen bearing species and its opacity becomes more significant compared to the competing absorption from H₂O and CH₄. This leads to dramatic changes in the spectrum. All the near infrared flux peaks are reduced by factors of 2-4. In particular, the new bands at short wavelengths dramatically affect the shape of the Y and J peaks at $T_{\rm eff} = 400 \, {\rm K}$. At the extremely low $T_{\rm eff}$ of 200 K, the changes are limited to the spectral regions were there are gaps in the HITRAN line list, and lead to very distinctive features in the Y and J bands. In particular, the Y band peak is split in two. Figure 6 suggests that in addition to the $NH_3 - H$ spectral index which more or less measures the slope of the blue side of the H band flux peak (Delorme et al. 2008), another NH₃-sensitive index could be defined to measure the curvature of the J band peak. In the mid-infrared (not shown), the most variation is shown by the 800 K spectrum with a < 40% decrease in flux in the $9-15\,\mu\mathrm{m}$ region. The 400 and 200 K spectra are barely affected in the mid-infrared. All these differences decrease somewhat as gravity is increased. The spectra shown in Fig. 6 and discussed above are computed in chemical equilibrium. However, we expect that the NH₃ abundance will be prevented from coming into equilibrium by convection, resulting in surface abundances ~ 10 times lower in the upper atmosphere. Consequently, NH₃ absorption is more muted in non-equilibrium spectra and the changes discussed above are smaller than in model that are in chemical equilibrium. The one exception is the $T_{\mathrm{eff}} = 200\,\mathrm{K}$ model which is so cold that the entire atmosphere is dominated by NH₃ whose abundance becomes nearly insensitive to convective mixing.

The recently identified Y dwarfs have $T_{\rm eff} \sim 300-500\,{\rm K}$ (Cushing et al. 2011). The spectra computed with the new BYTe line list indicate that broad NH₃ features should be detectable in the near infrared, most notably as a shoulder on the blue side of the Y band peak at $T_{\rm eff} \sim 400\,{\rm K}$ which is split at $T_{\rm eff} \lesssim 300\,{\rm K}$ by a growing NH₃ band centered at 1.03 μ m. The J band peak becomes triangular and the H band peak narrows. The strength of these effects is reduced by the non-equilibrium abundance of NH₃.

4. Models including both the new H₂ CIA and NH₃ opacities

We have computed new model atmospheres and synthetic spectra using the new H₂ CIA and NH₃ opacities. In all other aspects, these models are identical to those described in Saumon & Marley (2008); Marley et al. (2002) with additional details provided in Cushing et al. (2008) and Stephens et al. (2009). These new models, which include cloudless and cloudy atmospheres, were used in the recent analyzes presented in Cushing et al. (2011); Luhman et al. (2011) and Leggett et al. (2011). These new cloudless model spectra are compared to our previous models in Fig. 7, where, in contrast with Figs. 3 and 6, the (T, P) structures are computed with their respective opacities and thus the model structures and spectra are self-consistent. In this case, the flux is redistributed along the SED to conserve the total flux (which was not the case in Figs. 3 and 6). At $T_{\rm eff} = 1500$ and $1000 \, \rm K$, the changes are modest and dominated by the new H₂ CIA opacity. The reduced opacity in the first overtone band of H_2 - H_2 CIA is responsible for the increased flux in the K band peak ($\sim 15\%$) and a smaller increase in the H band flux. The $T_{\rm eff} = 1500\,\rm K$ spectrum also shows a 7% flux increase in the $4 \mu m$ peak but the 3-15 μm cloudless spectra are otherwise unaffected. Near the T/Y dwarf transition, where $T_{\rm eff} \sim 500 \, {\rm K}$, all four near infrared peaks in the spectrum are noticeably affected. Absorption by NH_3 changes the shape of the Y and J peaks and shaves off the blue side of the H band peak. Back warming from this NH_3 absorption pushes the flux out in the K band, which is now $\sim 75\%$ brighter, and in the $4-8 \,\mu\mathrm{m}$ region where the flux increases by $\sim 30\%$.

4.1. Comparison with the spectrum of Gliese 570D

The T7.5 dwarf Gliese 570D (Burgasser et al. 2000) is the most thoroughly studied late T dwarf. Optical, near infrared, M band, and Spitzer IRS spectroscopy are available – sampling $\sim 70\%$ of the total flux emitted – as well as Spitzer IRAC and WISE photometry. As this brown dwarf is a companion to a nearby main sequence star, its distance and metallicity are accurately known and its age is fairly well constrained to 2–5 Gyr. Because Gl 570D is nearby and relatively bright, the whole data set is of high quality and very effectively constrains the models. It is an excellent test object to validate the new models and evaluate the improvements brought about by the new opacities. Detailed analyzes of Gl 570D (Geballe et al. 2001; Saumon et al. 2006; Geballe et al. 2009) have established that $T_{\rm eff} = 800 - 820\,{\rm K}$ and $\log g = 5.09 - 5.23$ (cgs), where the ranges of $T_{\rm eff}$ and $\log g$ are correlated: a higher $T_{\rm eff}$ corresponds to a higher gravity. Solar metallicity was adopted based on the metallicity of the primary ($[Fe/H] = 0.09 \pm 0.04$). Saumon et al. (2006) established that convective transport drives the NH₃ abundance below the value expected from chemical equilibrium and obtained an very good fit of the entire SED of Gl 570D (see also Geballe et al. (2009)). Figure 8 shows the spectrum computed by Saumon et al. (2006) for those parameters within the above range that best reproduce the data and a new model computed with the same atmospheric parameters but with the new H₂ CIA and NH₃ opacities. There is significant improvement in the K band peak which is now fitted very well. There is also a modest improvement in the $5.5-7.5\,\mu\mathrm{m}$ region and beyond $9\,\mu\mathrm{m}$ where NH₃ absorption features overlap with H₂O absorption. The new H₂ CIA calculation, which is primarily responsible for the increased flux in the K band represents a significant improvement over the previously available tabulation. Outside of the 9–14 μ m region, NH₃ absorption is rather weak at this $T_{\rm eff}$, and the new NH₃ opacity leads to very modest changes. We believe that the remaining discrepancies in our fot of Gl 570D, such as the width of the J band peak and the depth of the $1.6\,\mu\mathrm{m}$ band of $\mathrm{CH_4}$, are primarily due to the very incomplete methane line list, which is presently in a similar state as the ammonia line list was prior to the first-principles calculation of Yurchenko, Barber & Tennyson (2011).

4.2. Comparison with near-infrared photometry

While a comparison with the SED of Gl 570D provides a detailed test of the new models, the overall effect of the new opacities can be better appreciated in color-magnitude

diagrams where synthetic colors can be compared to a large set of photometric data. Prior to 2003, our models used the solar abundances of Lodders & Fegley (1998) and the cloudless sequence was in generally good agreement with the near infrared colors of late T dwarfs (Burgasser et al. 2002). A significant downward revision of the CNO abundances in the solar photosphere (Allende Prieto et al. 2002) motivated a change of the solar abundances in our models (Lodders 2003) which moved the cloudless sequence to the blue in J-K. The resulting sequence of [M/H]=0 cloudless models are too blue compared to late field T dwarfs, particularly in J-K, when compared to the near infrared colors of late-T field dwarfs (Fig. 9 and Fig. 10e of Saumon & Marley (2008)) ³. The new H₂-H₂ CIA has a weaker first overtone band which increases the K band flux and moves the cloudless models to redder J-K, largely compensating for the effect of the reduced CNO abundances (Fig. 9). The new sequence of models is still slightly bluer by $\lesssim 0.15$ than the late T dwarf sequence which may be due to missing CH₄ opacity on the J band peak. The spread of the observed sequence of late T dwarf colors is significant and can be accounted for by the observed distribution in metallicity of field dwarfs (Fig. 15f of Saumon & Marley (2008)). The very latest T dwarfs recently discovered, with $M_K \gtrsim 17$ reveal a saturation in J-K and are redder than the calculated cloudless sequences. This may be caused by several limitations in our models such as the near infrared opacity of CH₄, the presence of clouds of compounds with low condensation temperature such as Na₂S, or the appearance of water clouds. Note that water condensation is included in the chemistry of the atmosphere and the gas phase H₂O abundance is reduced in the low- $T_{\rm eff}$ models where water condenses. We are investigating the role of clouds in low- $T_{\rm eff}$ atmospheres and the results will be reported in a separate publication. On the other hand, our cloudy sequences account for iron and silicate clouds and reproduce the SEDs and infrared colors of field L dwarfs fairly well (Saumon & Marley 2008; Cushing et al. 2008; Stephens et al. 2009). Cloudy models with $T_{\rm eff} \gtrsim 900\,{\rm K}$ are representative of field L dwarfs and the HR 8799 planets (Marley et al. 2012) and are shown in Fig. 9. These hotter, cloudy models are barely affected by the new opacities and their near infrared colors are very nearly identical to the older ones down to $T_{\rm eff} = 900 \, \rm K$. This is a consequence of the opacity of condensates which reduces the importance of the H₂ CIA and of the higher temperatures of the cloudy models which reduce the abundance of NH₃.

The same general picture emerges from the M_J versus J-H color-magnitude diagram (Fig. 10) although the new cloudless model now represents the J-H sequence of the late field T dwarfs quite well⁴. The data suggest that the J-H color saturates around -0.6 (see

³Note that the cloudless COND models of Allard et al. (2001) use the older solar abundances of Grevesse & Noels (1993) and agree very well with the observed near-infrared colors of late T dwarfs.

⁴This may be somewhat fortuitous in view of the model limitations mentioned above, particularly the

also Kirkpatrick et al. (2011) at $M_J \sim 17$ which is further supported by the position of the Y0 dwarf WISEP J1541 – 2250 in this color-magnitude diagram ($M_J = 23.9 \pm 0.8$, $J - H = 0.17 \pm 0.63$, Cushing et al. (2011); Kirkpatrick et al. (2011)). Our cloudless sequences do show such a minimum in J - H at $T_{\rm eff} \sim 400\,\rm K$ and a rapid turnover to the red at lower $T_{\rm eff}$. However, the models do not turn over to the red soon enough and reach values of $J - H \sim -1$, which is bluer than is observed. It will be interesting to see how the near infrared color-magnitude diagrams will be affected by a more complete CH₄ line list when it becomes available, or by the inclusion of a water cloud in the models.

5. Ammonia and Y dwarfs

Increased absorption in the near infrared has been observed on the blue side of the H band peak in the very late T dwarfs and in the new Y dwarfs and it is thought to be due to NH₃. The strength of the 1.5 μ m band of ammonia can be quantified with the NH₃ – H index defined by Delorme et al. (2008)

$$NH_3 - H = \frac{\int_{1.53}^{1.56} f(\lambda) d\lambda}{\int_{1.57}^{1.60} f(\lambda) d\lambda}.$$
 (2)

Smaller values of NH₃ – H indicate a relatively low flux on the blue side of the H band peak that is expected from stronger NH₃ absorption. A compilation of the data for the latest dwarfs known is shown in Fig. 11. The NH₃ – H index strongly correlates with the width of the J band peak (measured by the W_J index, Warren et al. (2007)), and with spectral type (Burningham et al. 2008; Cushing et al. 2011), with later types having smaller NH₃ – H. Figure 11 shows a break in the slope at $W_J \sim 0.3$ corresponding to spectral type T8 (Cushing et al. 2011). Based on the appearance of the blue side of the H band peak of the Y0 dwarf WISEP J1738+2732 – one of the most extreme objects in Fig. 11 – compared to earlier T8 and T9 dwarfs, Cushing et al. (2011) tentatively ascribe the absorption to NH₃.

The models shown in Fig. 11 reproduce the observed trend very well and indicate that the reduced abundance of NH₃ caused by non-equilibrium chemistry is favored. The dashed lines show the indices computed from the same models after the opacity of NH₃ has been removed with a trend that is strongly at odds with the data. The decrease in the modeled NH₃ – H at $T_{\rm eff} \lesssim 600\,{\rm K}$ (or spectral type later than T8) is caused by the 1.5 μ m band of NH₃. The expectation that NH₃ bands would appear in the near infrared at low enough $T_{\rm eff}$ and its detection as a depression in the blue side of the H band peak is thus well supported

poorly modeled $1.6 \,\mu\mathrm{m}$ band of CH₄ which affects the H band (Fig. 8)

on a quantitative basis by the models. However, the non-equilibrium models still show a systematic offset from the data, even for the low gravity of $\log g = 4$ (cgs) which, for these low $T_{\rm eff}$, would correspond to a rather low mass of $\sim 10\,{\rm M_{Jupiter}}$. It is very likely that the incomplete CH₄ line list is responsible for this systematic shift since methane is a strong absorber in the H band and a moderate absorber in the J band (Fig. 8).

Without the benefit of models, or with models of limited fidelity, the interpretation of the observed trends in spectral indices can be ambiguous, particularly when considering their variations over a wide range of T_{eff} . On the other hand, the direct detection of spectral features provides much more secure evidence of the presence of specific molecular species. While the detection of NH₃ in the $2.02-2.045\,\mu\mathrm{m}$ region of the spectrum of the T7p Gl 229B (Saumon et al. 2000) was tentative, Bochanski et al. (2011) identified 11 ammonia features in their medium resolution ($R \sim 6000$) near infrared spectrum of the T9 dwarf UGPS 072227.51-054031.2 (here after UGPS 0722-05). Figure 12 shows the portion of their spectrum with the most NH₃ features. For comparison, the top curve shows the log of the NH₃ opacity computed from the BYTe line list at the conditions corresponding to the atmospheric level where this part of the spectrum is emitted. The lowest curve is an unscaled model spectrum with $T_{\text{eff}} = 500 \,\text{K}$ and $\log g = 4.25$, solar metallicity, and $K_{zz} = 10^4 \,\text{cm}^2/\text{s}$ (Leggett et al. 2011). The six NH₃ absorption features identified by Bochanski et al. (2011) in this part of the spectrum all have corresponding troughs in the model spectrum. Four of these features also match peaks in the NH₃ opacity but the other two $(1.5264 \text{ and } 1.5411 \,\mu\text{m})$ do not and their attribution to NH₃ is not secure.

To better identify NH₃ absorption features in the spectrum of UGPS 0722-05 we show in Fig. 12 a spectrum computed from the same atmosphere model but without any NH₃ opacity (blue curve). There is obviously much NH₃ absorption over the entire range shown, even considering the fact that the model spectrum has a reduced (non-equilibrium) abundance of NH₃. This is fully consistent with the smaller value of the NH₃ – H index of UGPS 0722-05 compared to earlier T dwarfs (Fig. 11). However, a careful comparison of the spectra with and without NH₃ does not reveal any distinct feature at this spectral resolution. Every feature in one spectrum appears in the other, albeit with a different amplitude. For example, the 1.5142 μ m absorption feature matches a peak in the NH₃ opacity and an edge in the model spectrum (Fig. 12), but that edge also appears in the spectrum computed without NH₃, making the attribution doubtful. The model spectrum reproduces individual features of the observed spectrum fairly well (after scaling), but not exactly, which makes the attribution of specific features problematic. We find this to be also the case in other regions where Bochanski et al. (2011) found NH₃ features (1.21-1.27 μ m, 1.56-1.64 μ m and

at $1.990 \, \mu \text{m}$).⁵

The situation is different in the 1.96–2.05 μ m spectral region where Saumon et al. (2000) found 3 ammonia features in Gl 229B. Figure 13 shows this spectral region with the same models and opacity shown in Fig. 12 (with different scalings for clarity). The vertical solid and dashed lines correspond to 19 NH₃ absorption features that are clearly identifiable from the difference between the spectrum computed with the non-equilibrium NH₃ abundance (red) and the spectrum computed without NH₃ opacity (blue). In this spectral range, almost every peak in the NH₃ opacity (green) corresponds to a distinct absorption feature in the synthetic spectrum. The solid lines identify 8 absorption features that are present in the spectrum of UGPS 0722-05⁶, and the dashed lines indicate 6 NH₃ features that are not clearly detected in the spectrum and 5 that appear to be missing. Two of the detected features $(2.0374 \text{ and } 2.0455 \,\mu\text{m})$ match those found in Gl 229B by Saumon et al. (2000). An absorption feature clearly identified in Bochanski et al. (2011) at $1.9905 \,\mu\mathrm{m}$ (dotted line) corresponds to a well-defined absorption feature in the model spectrum. This feature is barely affected by the removal of all NH₃ opacity, however, and does not match very well with the NH₃ opacity peak centered at $1.9896 \,\mu\mathrm{m}$. In our model spectrum, this absorption is caused by H_2O .

We conclude that the detection of ammonia absorption features in the near infrared spectra of very late T dwarfs and Y dwarfs is easier in the K band near $2\,\mu\mathrm{m}$ than in the H band near $1.53\,\mu\mathrm{m}$. The main reason for this is that the NH₃ opacity in the K band is composed of a series of more or less equally spaced and well separated peaks (Fig. 13) while in the H band it has a denser irregular structure with a smaller dynamic range as well as a peak opacity that is about one order of magnitude smaller (Figs. 12 and 4). With higher resolving power it becomes possible to isolate NH₃ features in the H band. For example, with $R \sim 20000$, about a dozen features stand out in the 1.51–1.57 $\mu\mathrm{m}$ region of the model shown in Fig. 12. For dwarfs cooler than $\sim 400\,\mathrm{K}$, the condensation of water becomes significant (Burrows et al. 2003), removing H₂O from the gas and unmasking the NH₃ bands. For instance, the blue side of the H band peak becomes completely dominated by NH₃ absorption and NH₃ features should be readily detectable in the spectrum. This is also true in the $2-2.05\,\mu\mathrm{m}$ region, although the flux is much lower than at 1.55 $\mu\mathrm{m}$. The

⁵For completeness, we note that the two features in the low-resolution ($R \sim 500$) spectrum of UGPS 0722-05 pointed out by Lucas et al. (2010) at $\sim 1.275 \,\mu\mathrm{m}$ and at $\sim 1.282 \,\mu\mathrm{m}$ can be identified with synthetic spectra at higher resolution as blends of H₂O features. A weaker blend of H₂S features also contributes to the latter.

⁶If we ignore the single high-valued pixel making up the spike in the data at $2.0175 \,\mu\text{m}$, the NH₃ feature stands out.

NH₃ bands will thus become stronger with lower $T_{\rm eff}$ until ammonia itself begins to condense at $T_{\rm eff} \sim 160\,{\rm K}$ (Burrows et al. 2003).

6. Conclusion

We have computed new cloudy and cloudless models and synthetic spectra⁷ of brown dwarf atmospheres using recent first-principles calculations of the opacity of NH₃ and of the collision-induced absorption of H₂. Both constitute very significant theoretical improvements in the modeling of the opacity and, in the case of NH₃, the BYTe line list greatly expands over the HITRAN line list that is based primarily on experimental results. The cloudy models, relevant to L dwarfs, are barely affected by the new opacity because of their higher $T_{\rm eff}$ where NH₃ has a very low abundance and of the condensate opacity that largely conceals the H₂ CIA ⁸. We have thus focused on the role of these two opacities in cloudless models, which are relevant to spectral types \sim T4 and later.

The most notable change in the collision-induced absorption is a reduction of the opacity in the first roto-vibrational band centered at $4200\,\mathrm{cm^{-1}}$ (2.4 $\mu\mathrm{m}$), resulting in an increased K band flux. This corrects a deficiency in our earlier fit of the SED of Gl 570D (Saumon et al. 2006) and we now obtain a remarkable fit of nearly the entire SED of this late T dwarf (Fig. 8). Another consequence is that the cloudless sequence shifts toward bluer colors in near infrared color-magnitude diagrams and is in much better agreement with the colors of the late T dwarfs in the field. This new accurate H_2 CIA opacity will strengthen the analysis of gravity and metallicity indicators in the spectra of late T dwarfs and L subdwarfs and help reduce the uncomfortably large uncertainty in spectroscopic determinations of the gravity of brown dwarfs.

The new BYTe line list for NH_3 greatly improves the near infrared opacity of ammonia by adding several new bands at short wavelengths as well as a very large number of "hot transitions" that fill in gaps in the HITRAN line list. We can now reliably calculate the opacity of NH_3 at temperatures of several hundreds of degrees, which is crucial even for the coolest known dwarfs as the Y, J and H bands typically probe temperatures well above T_{eff} . The importance of this line list for the study of the T/Y transition and the new Y dwarfs cannot be overstated. Our models show that NH_3 develops very distinct signatures in the Y and J band flux peaks, and confirms the tentative identification of NH_3 absorption

⁷The synthetic spectra are available upon request to D. Saumon (dsaumon@lanl.gov).

 $^{^{8}}$ This is no longer the case in low-metallicity L subdwarfs where H₂ CIA is noticeable (Burgasser et al. 2003a, 2009; Cushing et al. 2009).

on the blue side of the H band peak which had been anticipated by Burrows et al. (2003). More quantitatively, the new models reproduce the trend in the spectral indices $NH_3 - H$ and W_J , confirming that the rapid decrease in NH₃ – H at spectral types later than T8 is due to NH₃ absorption. Thus, all the new WISE objects assigned to the Y dwarf class by Cushing et al. (2011) show NH₃ absorption in the H band. Furthermore, the data favors models where vertical transport reduces the NH₃ abundance in the observable part of the atmosphere. At a more detailed level, the medium resolution near infrared spectrum of the T9 UGPS 0722-05 (Bochanski et al. 2011) provides a rare opportunity to compare data with the detailed predictions of spectral features by the models and of the underlying molecular opacities. While we found that none of the features pointed out by Bochanski et al. (2011) can be unambiguously attributed to NH₃, our new models allow the identification of 9 other NH_3 absorption features around $2 \mu m$. The models clearly show that while there is strong NH₃ absorption in the H band in late T dwarfs and in the Y0 dwarfs, the general behavior of the NH₃ absorption in the 1.55 μ m region makes it more difficult to isolate individual features than in the 2.0 μ m region, requiring a resolving power of the order of $R \sim 20000$ compared to ~ 6000 in the K band.

Despite the progress that we have demonstrated here, obvious problems remain with the models. For instance, the fit of Gl 570D still has discrepancies that stand out. The near infrared color-magnitude diagrams for field dwarfs fail to reproduce the saturation (and possible turnover) revealed by the data at very low $T_{\rm eff}$. The spectral indices NH₃ – H and W_J do not yet provide a quantitative diagnostic of the atmospheric structure and chemistry. With these two new calculations of molecular opacities that affect primarily late T dwarfs and the new class of Y dwarfs, reliable opacities are available for all important gas phase absorbers except for CH₄. The inadequacy of the the current line list of methane has been much maligned ever since the very first spectral modeling of a T dwarf (Marley et al. 1996; Allard et al. 1996). We believe that for cloudless atmosphere models, the CH₄ opacity is the only remaining obstacle of significance to modeling the spectroscopic data accurately. The availability of a CH₄ line list comparable in quality and scope to the BYTe NH₃ line list would open the door to reliable determinations of T dwarf gravities and metallicities, two fundamental parameters that have remained elusive for the vast majority of brown dwarfs.

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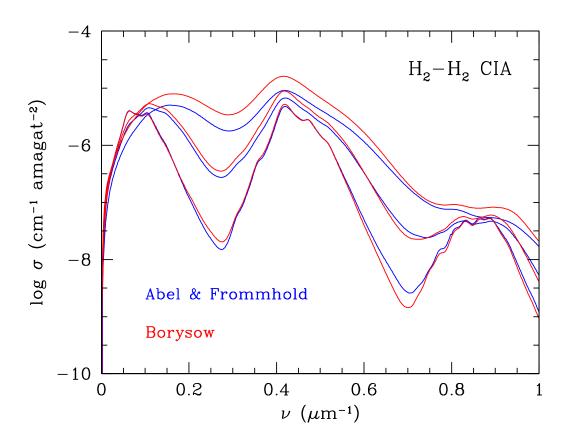


Fig. 1.— Collision-induced absorption coefficient for H_2 - H_2 collisions at T=500, 1000 and 2000 K, from bottom to top, respectively. Each broad peak corresponds to a change in the vibrational quantum number of $\Delta v=0$, 1, 2... from left to right, respectively. The calculations of Borysow and collaborators (Borysow et al. (2001); Borysow (2002), and references therein) and the recent work of Abel & Frommhold (Frommhold et al. 2010; Abel et al. 2011) are shown in red and blue, respectively. 1 amagat= $2.6867774 \times 10^{19} \,\mathrm{cm}^{-3}$. [See the electronic edition of the Journal for a color version of this figure.]

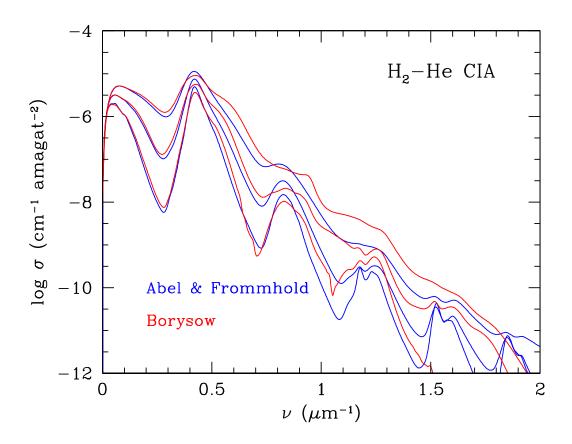


Fig. 2.— Same as Figure 1 for H_2 -He CIA. The older calculation labeled "Borysow" is from Borysow et al. (1989), Jorgensen et al. (2000), and references therein. The "Abel & Frommhold" calculation is from Abel et al. (2012). [See the electronic edition of the Journal for a color version of this figure.]

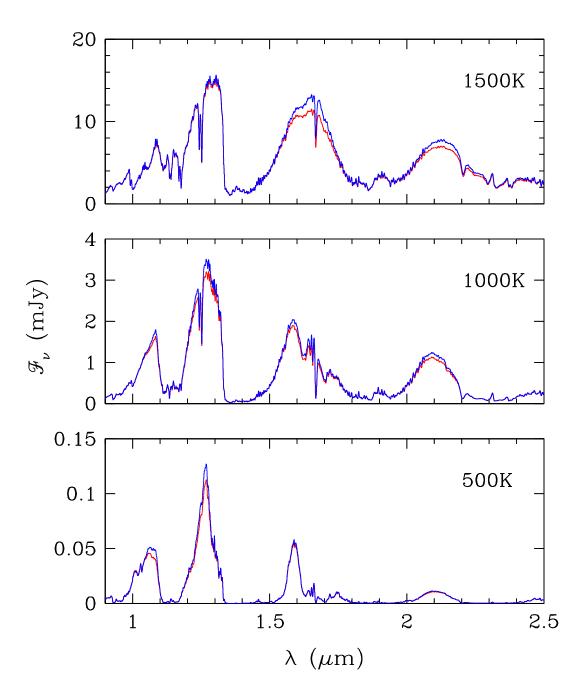


Fig. 3.— Effect of the new H_2 - H_2 and H_2 -He CIA opacity on synthetic spectra of brown dwarfs. The spectra shown are *cloudless* models with $T_{\rm eff}=1500,\,1000$ and $500\,\rm K$, with $\log g=5$ (cgs) and solar metallicity. The spectra computed with the new CIA opacities are shown in blue. The red lines show spectra computed with the older CIA opacity and the same (T,P) structures. The fluxes are calculated for $d=10\,\rm pc$ and are displayed at a resolving power R=500.

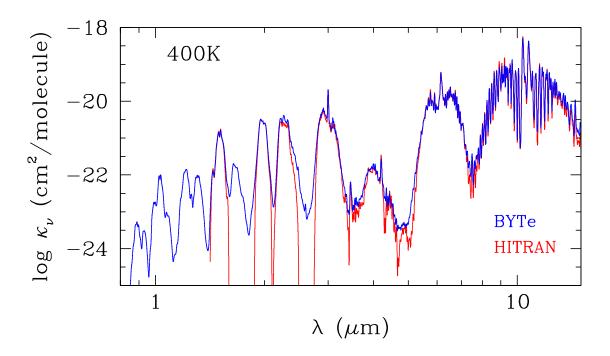


Fig. 4.— Absorption coefficient of NH₃ at $T=400\,\mathrm{K}$ and $P=1\,\mathrm{bar}$. The red line shows the opacity calculated from the line list that we have used in our previous atmosphere models. It is primarily based on the HITRAN data base supplemented with more recent laboratory measurements (Freedman et al. 2008). The blue line labeled "BYTe" shows the absorption coefficient computed from the new first-principles line list of Yurchenko, Barber & Tennyson (2011). For clarity, the absorption coefficient is shown at a resolving power of R=200.

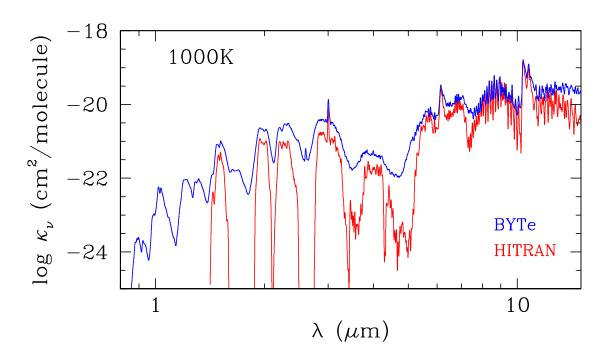


Fig. 5.— Same as Figure 4 for $T=1000\,\mathrm{K}.$

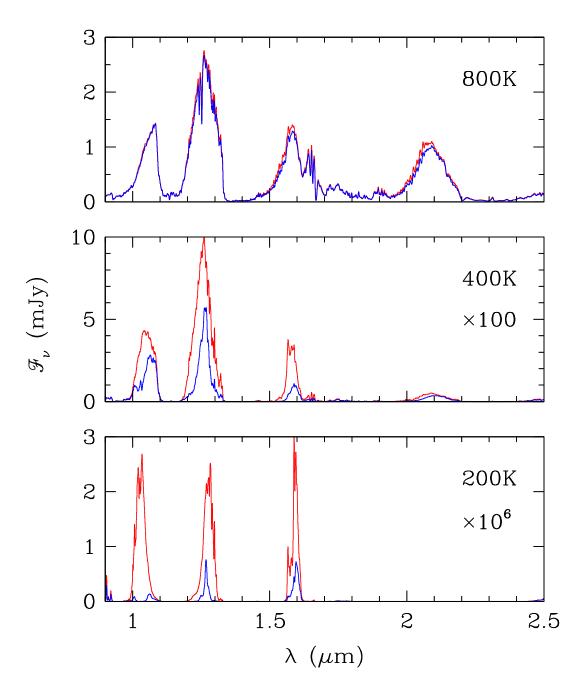


Fig. 6.— Effect of the new NH₃ opacity on near infrared spectra of cool brown dwarfs. Three models are shown with $T_{\rm eff}=800,\,400$ and 200 K. All models shown have solar composition, log g=4, are cloudless and in chemical equilibrium. Models computed with the HITRAN line list for NH₃ are shown in red while those computed from the same (T,P) structure but with the BYTe NH₃ line list are in blue. The spectra are shown as absolute fluxes for a distance of 10 pc and at R=500 for clarity.

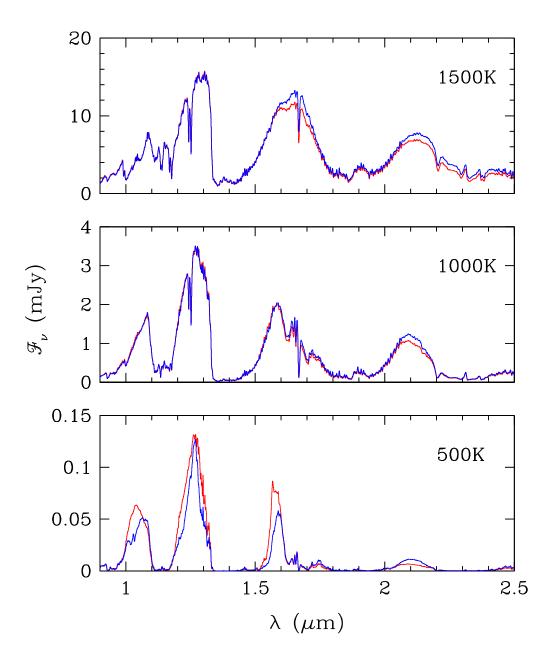


Fig. 7.— Comparison of spectra including the new H_2 CIA and the BYTe NH_3 opacities (blue) with spectra from a previous generation that only differ in these two sources of opacity (red). These cloudless spectra have $\log g = 5$ (cgs), solar metallicity and are computed in chemical equilibrium. The effective temperatures are 1500, 1000 and 500 K. In contrast with Figs. 3 and 6, the atmospheric (T, P) structures have been computed with the appropriate (old or new) opacities. The spectra are plotted at R = 500 for clarity.

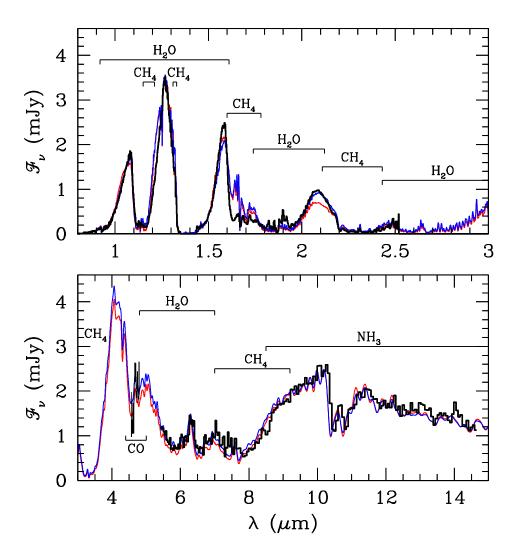


Fig. 8.— Spectrum of Gliese 570 D. The data are shown in black (Burgasser et al. 2003b; Geballe et al. 2001; Cushing et al. 2006; Geballe et al. 2009). The spectrum obtained with our previous models is shown in red (Saumon et al. 2006). The blue curve shows the spectrum computed with the new H_2 CIA and the BYTe NH₃ line list with the same model parameters: $T_{\rm eff} = 820 \, {\rm K}$, $\log g = 5.23 \, {\rm (cgs)}$, $[{\rm M/H}]{=}0$, and eddy diffusion coefficient $\log K_{zz} \, {\rm (cm^2/s)}{=}4.5$. The models are plotted at a resolving power of R = 500 (top panel) and R = 100 (bottom panel), approximating that of the data. The models are not normalized to the data but are fluxes at Earth computed with the known distance of Gl 570D and the radius obtained from the evolution with the above parameters.

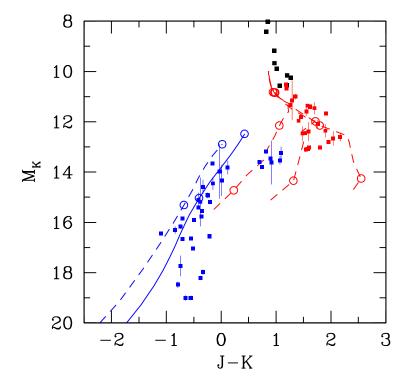


Fig. 9.— M_K versus J-K color-color diagram (MKO system) comparing models with the near infrared photometry of field MLT dwarfs. The photometry is from Leggett et al. (2002), Knapp et al. (2004), and Marocco et al. (2010) with M dwarfs in black, L dwarfs in red and T dwarfs in blue. All known binaries have been removed from the sample except those with resolved MKO photometry: ϵ Indi B (McCaughrean et al. 2004), SDSS J102109.69-030420.1 and SDSS J042348.57-041403.5 (Burgasser et al. 2006), and Kelu-1 (Liu & Leggett 2005). The parallaxes are from Perryman et al. (1997), Dahn et al. (2002), Tinney et al. (2003), Vrba et al. (2004), Marocco et al. (2010), and various references in Leggett et al. (2002). Solid blue curve: Cloudless models calculated with the new H₂ CIA and the BYTe NH₃ line list and an eddy diffusion coefficient $K_{zz} = 10^4 \,\mathrm{cm}^2/\mathrm{s}$ (e.g. these model are not in chemical equilibrium). The curve extends from $T_{\rm eff}=1500\,{\rm K}$ to $500\,{\rm K},$ with open circles indicating $T_{\rm eff} = 1500\,{\rm K}$ and $1000\,{\rm K}$. The dashed blue curve shows models computed with older ${\rm H_2}$ CIA and NH₃ opacities and $K_{zz} = 0$ (chemical equilibrium) (Saumon & Marley 2008). The dashed red curves are cloudy models in chemical equilibrium from Saumon & Marley (2008) extending from 2400 K to 900 K with open circles indicating (from the top) $T_{\rm eff} = 2000$, 1500, and 1000 K. Each curve corresponds to a different value of the cloud sedimentation parameter (see Ackerman & Marley (2001); Saumon & Marley (2008)) of $f_{\text{sed}} = 1$, 2 and 3, from right to left, respectively. Cloudy models computed with the new opacities overlap the cloudy sequences shown. All models shown have solar metallicity and $\log g = 5$.

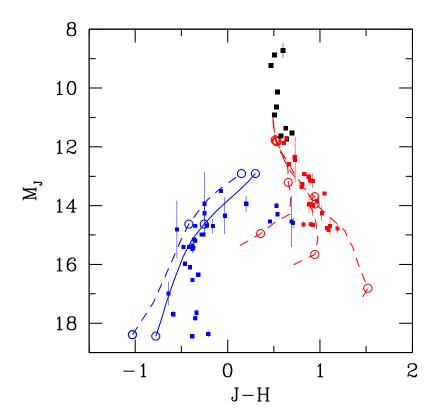


Fig. 10.— Same as Figure 9 for M_J versus J-H. The Y0 dwarf WISEP J1541-2250, with $M_J=23.9\pm0.8$ and $J-H=0.17\pm0.63$ (Cushing et al. 2011; Kirkpatrick et al. 2011), falls well outside this figure.

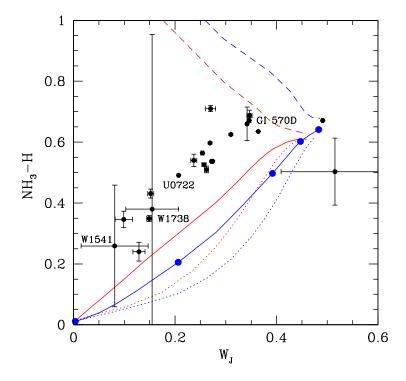


Fig. 11.— Spectral indices showing the growth of absorption in the blue wing of the H band of late T dwarfs. The spectral indices $NH_3 - H$ and W_J are defined in Delorme et al. (2008) and Warren et al. (2007), respectively. The curves show the indices computed from cloudless synthetic spectra starting at $T_{\rm eff} = 200$ (lower left corner) to $1000\,\rm K$ (right) with solid dots marking steps of 200 K. Two gravities, $\log g = 4$ (cgs, red curves) and 5 (blue), are shown. Dotted lines show models in chemical equilibrium and solid curves are models driven out of equilibrium by convective transport and mixing in the radiative zone parametrized by the eddy diffusion coefficient $K_{zz} = 10^4\,\rm cm^2/s$. The dashed lines show spectra computed with the same (T,P) structures after removing the NH₃ opacity. The dwarfs Gl 570D (T8, Delorme et al. (2008)), UGPS 0722-05 (T9, Cushing et al. (2011)), WISEP J1738+2732 (Y0, Cushing et al. (2011), and WISEP J1541-2250 (Y0, Cushing et al. (2011)) are labeled. Data compiled from Warren et al. (2007); Delorme et al. (2008); Burningham et al. (2008, 2009, 2011a,b); Cushing et al. (2011), and Liu et al. (2011).

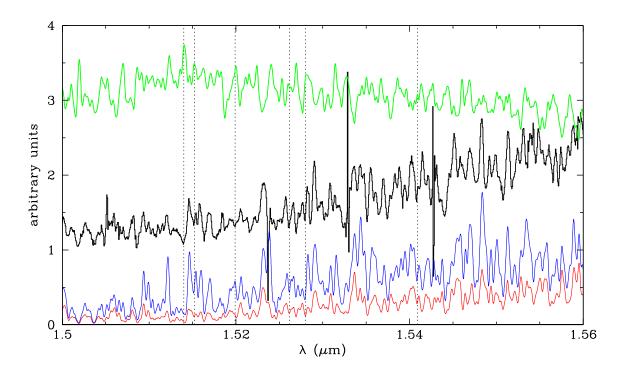


Fig. 12.— Medium resolution spectrum ($R \sim 6000$) of UGPS0722-05 (Bochanski et al. 2011) (black). The observed spectrum has been rescaled, shifted vertically, and the radial velocity (47 km/s, Bochanski et al. (2011)) has been removed. A cloudless synthetic spectrum, with $T_{\rm eff} = 500 \, {\rm K}$, $\log g = 4.25$, [M/H]=0, $K_{zz} = 10^4 \, {\rm cm}^2/{\rm s}$ (Leggett et al. 2011) is shown in red. The blue curve represents the same spectrum calculated without any NH₃ opacity. The \log_{10} of the NH₃ opacity at 650 K and 1 bar, corresponding to the emission level of this part of the spectrum in the model, is shown in green at an arbitrary scale. The model spectra and the opacity are shown at the spectral resolution of the data. The vertical dotted lines show the NH₃ absorption features identified by Bochanski et al. (2011) in the spectrum of UGPS 0722-05 which should correspond to peaks in the NH₃ opacity.

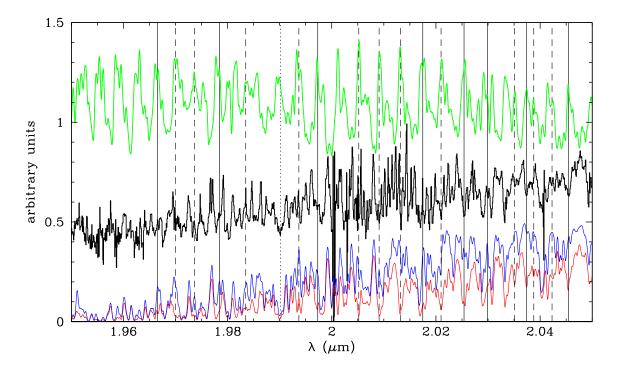


Fig. 13.— Same as Fig. 12 but in the K band. The models have been rescaled for clarity. The vertical dotted line shows the NH₃ feature identified by Bochanski et al. (2011) in the spectrum of UGPS 0722-05. The other vertical lines are NH₃ absorption features predicted by the model spectrum, which all match peaks in the NH₃ opacity. Those that are also apparent in the data are shown with solid lines while those that are ambiguous or missing are shown with dashed lines.